FDA 510(k) Premarket Notification

# K781932 BLOOD PRESSURE RECORDER

**CLEARED**

Nov 29, 1978
9 days to decision

K781932 · Product code: **DXN** · Cardiovascular
Source: https://www.510kdatabase.net/k781932/

## SUBMISSION DETAILS

| | |
|---|---|
| Decision | Substantially Equivalent (Cleared) |
| Submission type | Traditional |
| Device classification | System, Measurement, Blood-pressure, Non-invasive (DXN) |
| Date received | Nov 20, 1978 |
| Decision date | Nov 29, 1978 |
| Days to decision | 9 days |
| Third-party review | No |

## APPLICANT

| | |
|---|---|
| Company | **Cardiodyne, Inc.** |
| Location | Mchenry, IL, US |
| 510(k) history | 3 submissions · 3 cleared · 1977-1991 |

**510k Database** - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k781932/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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